CLAIMS

1. A compound of formula I:

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wherein R^4 is selected from:

$$-CH=C_{\mathcal{M}}^{\mathcal{X}} \xrightarrow{(O)_{m}} \text{ and } -CH=C_{\mathcal{M}}^{\mathcal{X}} CO-Z$$

X represents H, halogen, CN or methyl;

R¹ represents H or C₁₋₄alkyl which is optionally substituted with OH or C₁₋₄alkoxy; or R¹ and R² together complete a heterocyclic ring of 3-7 members bearing 0-2 substituents, in addition to R³, selected from halogen, oxo, NO₂, CN, CF₃, C₁₋₆alkyl, C₂₋₆acyl, C₂₋₆alkenyl, C₁₋₆alkoxy, C₁₋₆alkoxycarbonyl and Ar;

when R¹ represents H or optionally substituted C₁-₄alkyl, R² and R³ independently represent H, C₁-₁₀alkyl, C₃-₁₀cycloalkyl, C₃-₆cycloalkylC₁-₆alkyl, C₂-₁₀alkenyl, C₂-₁₀alkynyl, Ar, heterocyclyl, or heterocyclylC₁-₆alkyl, wherein the alkyl, cycloalkyl, alkenyl and alkynyl groups optionally bear one substituent selected from halogen, CF₃, NO₂, CN, Ar, ArCH₂O, ArO, -OR¹¹, -SR¹¹, -SO₂R¹², -COR¹¹, -CO₂R¹¹, -CON(R¹¹)₂, -OCOR¹², -N(R¹¹)₂ and -NR¹¹COR¹²; and the heterocyclic groups optionally bear up to 3 substituents independently selected from halogen, NO₂, CN, R¹², Ar, ArCH₂O, ArO, ArOCH₂, -OR¹¹, -SR¹¹, -SO₂R¹², -COR¹¹, -CO₂R¹¹, -CO₂R¹¹, -CON(R¹¹)₂, -OCOR¹², -N(R¹¹)₂ and -NR¹¹COR¹²;

or R² and R³ together with the nitrogen to which they are mutually attached complete a mono- or bicyclic heterocyclic ring system of 5-10 ring

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atoms selected from C, N, O and S, said ring system optionally having an additional benzene or heteroaryl ring fused thereto, said heterocyclic system and optional fused ring bearing 0-3 substituents independently selected from halogen, oxo, NO₂, CN, R¹², Ar, ArCH₂O, ArO, ArOCH₂, -OR¹¹, -SR¹¹, -SO₂R¹², -COR¹¹, -CO₂R¹¹, -CON(R¹¹)₂, -OCOR¹², -N(R¹¹)₂ and -NR¹¹COR¹²;

and when R^1 completes a ring with R^2 , R^3 represents H, C_{1-6} alkyl, C_{2-6} acyl, C_{2-6} alkenyl or benzyl;

m is 0 or 1, with the proviso that when m is 1 neither R^2 nor R^3 is H and R^3 is not acyl, and that m is 1 when X and R^1 are both H;

R¹¹ represents H or R¹²;

R¹² represents C₁₋₆alkyl which optionally bears up to 3 halogen substituents or one substituent selected from CN, OH, C₁₋₄alkoxy and C₁₋₄alkoxycarbonyl;

Y represents halogen, CN or methyl;

Z represents OR11 or N(R5)R6;

 R^5 and R^6 have the same definition as R^2 and R^3 in the embodiment in which R^1 is H or optionally substituted C_{1-4} alkyl;

R¹⁴ represents H or C₁₋₆alkyl, C₃₋₇cycloalkyl, C₃₋₆cycloalkylC₁₋₆alkyl,
C₂₋₆alkenyl, C₂₋₆alkynyl, phenyl or benzyl, any of which optionally bear up
to 3 halogen substituents or one substituent selected from CN, NO₂, OH,
C₁₋₄alkoxy, CO₂H, C₁₋₄alkoxycarbonyl, C₂₋₆acyl, C₂₋₆acyloxy, amino,
C₁₋₄alkylamino, di(C₁₋₄alkyl)amino, C₂₋₆acylamino, carbamoyl,
C₁₋₄alkylcarbamoyl and di(C₁₋₄alkyl)carbamoyl; and

Ar represents phenyl or heteroaryl either of which optionally bears up to 3 substituents independently selected from halogen, CF₃, NO₂, CN, OCF₃, C₁₋₆alkyl and C₁₋₆alkoxy; or a pharmaceutically acceptable salt thereof.

30 2. A compound according to claim 1 of formula II:

or a pharmaceutically acceptable salt thereof.

- 5 3. A compound according to claim 2 wherein R¹ and R² complete a heterocyclic ring of 5 or 6 atoms and R³ represents H; C¹-6alkyl, C²-6acyl or benzyl.
- 4. A compound according to claim 2 wherein R¹ is H or optionally
 10 substituted C¹-4alkyl and R² and R³ complete a heterocyclic ring system.
 - 5. A compound according to claim 4 wherein R^{14} is 2,2,2-trifluoroethyl, X is F, CN or methyl, and R^1 is H.
- 15 6. A compound according to claim 4 wherein m is 1 and X and R^1 are both H.
 - 7. A compound according to claim 1 of formula III:

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or a pharmaceutically acceptable salt thereof.

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- 8. A compound according to claim 7 wherein Y represents F, CN or methyl and Z represents OH, C₁₋₆alkoxy or N(R⁵)R⁶.
- 9. A compound according to claim 8 wherein R¹⁴ represents 2,2,2-5 trifluoroethyl and Z represents ethoxy.
 - 10. A pharmaceutical composition comprising a compound according to any previous claim and a pharmaceutical carrier.
- 10 11. A compound according to any of claims 1-9 for use in a method of treatment of the human body.
 - 12. The use of a compound according to any of claims 1-9 in the manufacture of a medicament for treating or preventing Alzheimer's disease.

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13. A method of treatment of a subject suffering from or prone to Alzheimer's disease comprising administering to that subject an effective amount of a compound according to any of claims 1-9.